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                 STN AnaVist, Version 2.0, now available with Derwent
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                 FORIS renamed to SOFIS
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                 CAplus coverage extended to include traditional medicine
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         SEP 17
                 patents
                 EMBASE, EMBAL, and LEMBASE reloaded with enhancements
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         SEP 24
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NEWS 23
         OCT 02
                 Zentralblatt
NEWS 24
         OCT 19
                 BEILSTEIN updated with new compounds
NEWS EXPRESS 19 SEPTEMBER 2007: CURRENT WINDOWS VERSION IS V8.2,
              CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0jc(jp),
              AND CURRENT DISCOVER FILE IS DATED 19 SEPTEMBER 2007.
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=> s primase and rna and (fluorescen? or fluorophore) and (template or target) 19 PRIMASE AND RNA AND (FLUORESCEN? OR FLUOROPHORE) AND (TEMPLATE L1OR TARGET)

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4 DUP REMOVE L4 (0 DUPLICATES REMOVED)

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ANSWER 1 OF 4 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN L5ΤT High throughput screening assays for bacterial primases

L5ANSWER 2 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

TI Fluorometric assay for bacterial primases

L5ANSWER 3 OF 4 MEDLINE on STN

TI Homogenous assays for Escherichia coli DnaB-stimulated DnaG primase and DnaB helicase and their use in screening for chemical inhibitors.

L5ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

TIFlashPlate scintillation proximity assays for characterization and screening of DNA polymerase, primase, and helicase activities

=> d bib 1-4

L5: ANSWER 1 OF 4 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN

- AN 2006:345752 BIOSIS
- DN PREV200600344884
- TI High throughput screening assays for bacterial primases
- AU Griep, Mark A. [Reprint Author]; Koepsell, Scott A.; Hinrichs, Steven H.
- CS Univ Nebraska, Lincoln, NE 68588 USA
- SO FASEB Journal, (MAR 6 2006) Vol. 20, No. 4, Part 1, pp. A510-A511.

 Meeting Info.: Experimental Biology 2006 Meeting. San Francisco, CA, USA.

 April 01 -05, 2006. Amer Assoc Anatomists; Amer Physiol Soc; Amer Soc

 Biochem & Mol Biol; Amer Soc Investigat Pathol; Amer Soc Nutr; Amer Soc

 Pharmacol & Expt Therapeut.

 CODEN: FAJOEC. ISSN: 0892-6638.
- DT Conference; (Meeting)
 Conference; Abstract; (Meeting Abstract)
- LA English
- ED Entered STN: 12 Jul 2006 Last Updated on STN: 12 Jul 2006
- L5 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN
- AN 2005:265750 CAPLUS
- DN 142:477808
- TI Fluorometric assay for bacterial primases
- AU Koepsell, Scott A.; Hanson, Sarah; Hinrichs, Steven H.; Griep, Mark A.
- CS Department of Microbiology and Pathology, University of Nebraska Medical Center, Omaha, NE, 68198, USA
- SO Analytical Biochemistry (2005), 339(2), 353-355 CODEN: ANBCA2; ISSN: 0003-2697
- PB Elsevier
- DT Journal
- LA English
- RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT
- L5 ANSWER 3 OF 4 MEDLINE on STN
- AN 2002270850 MEDLINE
- DN PubMed ID: 12009693
- TI Homogenous assays for Escherichia coli DnaB-stimulated DnaG primase and DnaB helicase and their use in screening for chemical inhibitors.
- AU Zhang Yi; Yang Fude; Kao Yeh-Chih; Kurilla Michael G; Pompliano David L; Dicker Ira B
- CS Pharmaceutical Research Institute, Bristol-Myers Squibb Company, Wilmington, DE 19880, USA.
- SO Analytical biochemistry, (2002 May 15) Vol. 304, No. 2, pp. 174-9. Journal code: 0370535. ISSN: 0003-2697.
- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 200210
- ED Entered STN: 16 May 2002 Last Updated on STN: 11 Oct 2002 Entered Medline: 10 Oct 2002
- L5 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN
- AN 2001:154887 CAPLUS
- DN 134:337441
- TI FlashPlate scintillation proximity assays for characterization and screening of DNA polymerase, primase, and helicase activities
- AU Earnshaw, David L.; Pope, Andrew J.
- CS Molecular Interactions and New Assay Technologies, SmithKline Beecham Pharmaceuticals, Essex, UK
- SO Journal of Biomolecular Screening (2001), 6(1), 39-46

CODEN: JBISF3; ISSN: 1087-0571

- PB Mary Ann Liebert, Inc.
- DT Journal
- LA English
- RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT
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- L6 4 L1 AND SCREEN?
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- L6 ANSWER 1 OF 4 MEDLINE on STN
- TI Homogenous assays for Escherichia coli DnaB-stimulated DnaG primase and DnaB helicase and their use in screening for chemical inhibitors.
- L6 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Fluorometric assay for bacterial primases
- L6 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN
- TI FlashPlate scintillation proximity assays for characterization and screening of DNA polymerase, primase, and helicase activities
- L6 ANSWER 4 OF 4 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
- TI High throughput screening assays for bacterial primases
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- L7 ANSWER 1 OF 16 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
- TI High throughput screening assays for bacterial primases.
- L7 ANSWER 2 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Methods for real-time recombinase-polymerase amplification (RPA) of target DNA
- L7 ANSWER 3 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Methods and materials for RPA (recombinase polymerase amplification) of double stranded nucleic acids
- L7 ANSWER 4 OF 16 EMBASE COPYRIGHT (c) 2007 Elsevier B.V. All rights reserved on STN
- TI Crosstalk between primase subunits can act to regulate primer synthesis in trans.
- L7 ANSWER 5 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Fluorometric assay for bacterial primases
- L7 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides
- L7 ANSWER 7 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- TI Oligonucleotide tagged nucleoside triphosphates (OTNTPs) for genetic analysis, and synthesis from reactive bifunctional amidites

- L7 ANSWER 8 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- TIGenotype analysis using RecA protein and recombinase polymerase amplification (RPA) for potential use in molecular diagnosis of disease or detection of pathogenic organisms
- L7ANSWER 9 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- TIDetection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension
- L7 ANSWER 10 OF 16 MEDLINE on STN DUPLICATE 1
- Mechanism and stoichiometry of interaction of DnaG primase with TIDnaB helicase of Escherichia coli in RNA primer synthesis.
- L7 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- TIIdentification of differentially expressed genes in pancreatic cancer cells using cDNA microarray
- L7 ANSWER 12 OF 16 MEDLINE on STN
- Homogenous assays for Escherichia coli DnaB-stimulated DnaG TIprimase and DnaB helicase and their use in screening for chemical inhibitors.
- L7 ANSWER 13 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- FlashPlate scintillation proximity assays for characterization and screening of DNA polymerase, primase, and helicase activities
- L7 ANSWER 14 OF 16 MEDLINE on STN
- Amplifications of DNA primase 1 (PRIM1) in human osteosarcoma. TI
- L7 ANSWER 15 OF 16 MEDLINE on STN
- Structural and functional studies of the rat mitochondrial single strand TIDNA binding protein P16.
- L7 ANSWER 16 OF 16 MEDLINE on STN
- Identification and subcellular localization of the polypeptide for chick DNA primase with a specific monoclonal antibody.
- => d bib kwic 6,9,10 17
- L7 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN
- 2004:220076 CAPLUS AN
- 140:248188 DN
- Detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides
- Hanna, Michelle M. IN
- PΑ
- SO U.S. Pat. Appl. Publ., 104 pp., Cont.-in-part of Appl. No. PCT/US02/34419. CODEN: USXXCO
- DTPatent
- English LA
- FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	US 2004054162	A1	20040318	US 2003-425037	20030429
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	WO 2003038042	A2	20030508	WO 2002-US34419	20021029
	WO 2003038042	A3	20040325		

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     JP 2006525022
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     US 2003-425037
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TI
     Detection of nucleic acid sequences by isothermal RNA
     polymerase-dependent primer extension using reporter group-labeled
AB
              analogs may be incorporated into nucleic acids. In one
     embodiment, the process generates multiple amplification products from the
     primer and target. The methods generally comprise using a
     nucleoside, a mononucleotide, an oligonucleotide, or a polynucleotide, or
     analog thereof, to initiate synthesis of an oligonucleotide product that
     is substantially complementary to a target site on the defined
     polynucleotide sequence; optionally using nucleotides or nucleotide
     analogs as oligonucleotide chain elongators or chain terminators to.
     transcriptional amplification nucleic acid RNA polymerase
ST
     nucleotide nucleoside analog; DNA methylation analysis transcriptional
     amplification
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (CDKN2A, anal. of methylation of; detection of nucleic acid sequences
        by isothermal RNA polymerase-dependent primer extension using
        reporter group-labeled nucleotides)
IT
     Genetic element
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        (CpG island, anal. of methylation of; detection of nucleic acid
        sequences by isothermal RNA polymerase-dependent primer
        extension using reporter group-labeled nucleotides)
TT
     Bacteriophage SP6
     Coliphage T7
     Enterobacteria phage T3
     Escherichia coli
        (RNA polymerase of; detection of nucleic acid sequences by
        isothermal RNA polymerase-dependent primer extension using
        reporter group-labeled nucleotides)
IT
     Feces
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(anal. of DNA methylation in; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Methylation RL: BSU (Biological study, unclassified); BIOL (Biological study) (anal. of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Nucleotides, analysis Purine nucleotides Pyrimidine nucleotides RL: ARU (Analytical role, unclassified); ANST (Analytical study) (analogs, reporter group containing, in transcriptional primer elongation; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Fluorescent dyes (as reporter groups; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) RNA RL: ANT (Analyte); ANST (Analytical study) (as template for amplification; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Diagnosis (cancer, anal. of DNA methylation in; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) RL: ANT (Analyte); ANST (Analytical study) (conjugates with oligonucleotides, detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Nucleic acid amplification (method) (detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Primers (nucleic acid) RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Mutation (detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Pathogen (diagnostic detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) DNA microarray technology Northern blot hybridization Nucleic acid hybridization Southern blot hybridization (for capture and anal. of amplification products; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) Antibodies and Immunoglobulins RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (for protein capture; detection of nucleic acid sequences by isothermal

RNA polymerase-dependent primer extension using reporter

group-labeled nucleotides)

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TIFluorescence resonance energy transfer (in detection of transcriptional primer extension; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT RL: MOA (Modifier or additive use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses) (in protein conjugation with oligonucleotides; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT (mol.; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT Transcription, genetic (nucleic acid amplification using; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT Deamination (of 5-methylcytosine, in detection of DNA methylation; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT Cyanine dyes (oligonucleotide conjugates, as reporters; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT Quantum dot devices (primer conjugates, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) ΙT Phycoerythrins RL: ARU (Analytical role, unclassified); ANST (Analytical study) (primer conjugates, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) TT Promoter (genetic element) RL: ARU (Analytical role, unclassified); BUU (Biological use, unclassified); ANST (Analytical study); BIOL (Biological study); USES (primers containing, for transcriptional amplification; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) TΤ Nucleoside analogs RL: ARU (Analytical role, unclassified); ANST (Analytical study) (reporter group containing, in transcriptional primer elongation; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT RL: ANT (Analyte); BSU (Biological study, unclassified); ANST (Analytical study); BIOL (Biological study) (specific detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT Nucleic acid amplification (method) (transcriptional; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT 120-73-0D, Purine, analogs 289-95-2D, Pyrimidine, analogs 29220-54-0 185971-89-5 291536-62-4 400051-23-2D, AlexaFluor 647, conjugates with 670257-80-4 670257-82-6 670257-84-8 670257-86-0 671225-92-6 671234-25-6 671234-26-7 671234-27-8 671234-28-9 RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (as reporter, incorporation into primer extension products; detection

of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT 951-78-0, Deoxyuridine RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (detection in DNA in anal. of DNA methylation; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) 554-01-8, 5-Methylcytosine IT RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (detection in DNA of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT 9012-90-2, DNA-dependent DNA polymerase 9014-24-8, DNA-dependent 9026-28-2, RNA-dependent RNA RNA polymerase polymerase 64885-96-7, Primase RL: CAT (Catalyst use); USES (Uses) (detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT 2382-65-2D, methylated RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) 671527-54-1 671527-55-2 IT671527-51-8 671527-52-9 671527-57-4 671527-58-5 671527-59-6 671527-60-9 671527-61-0 671527-63-2 671527-64-3 671527-65-4 RL: PRP (Properties) (unclaimed nucleotide sequence; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) IT 671527-53-0 671527-62-1 RL: PRP (Properties) (unclaimed sequence; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension using reporter group-labeled nucleotides) L7 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2007 ACS on STN ΑN 2003:356568 CAPLUS 138:363805 DN Detection of nucleic acid sequences by isothermal RNA TIpolymerase-dependent primer extension IN Hanna, Michelle M. PA Ribomed, Inc., USA; Ribomed Technologies, Inc. SO PCT Int. Appl., 183 pp. CODEN: PIXXD2 DT Patent LA English FAN.CNT 3 PATENT NO. KIND DATE APPLICATION NO. DATE _ - - ------______ WO 2003038042 A2 WO 2002-US34419 20030508 PI 20021029 WO 2003038042 Α3 20040325 AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,

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      Detection of nucleic acid sequences by isothermal RNA
      polymerase-dependent primer extension
      A method for detection of a target nucleic acid sequence by
      RNA polymerase-dependent elongation of a primer is described.
                                                                      The
      primer is elongated by the polymerase until the enzyme incorporates a
      blocked.
               . . results in extension product termination. The polymerase
      may then initiate extension of a new primer leading to amplification of
      the target sequence. The primer may include a promoter sequence
      suitable for the RNA polymerase or a fluorescent dyes
      as reporters. In one aspect, the invention provides a method for
      detecting a target protein, DNA or RNA by generating
      multiple detectable RNA oligoribonucleotides by abortive
      transcription. The method can be used for genotyping, mol. diagnosis, and
      detection of DNA methylation.
. ST
      transcriptional amplification nucleic acid RNA polymerase
      primer; DNA methylation analysis transcriptional amplification
IT
      Gene, animal
      RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (CDKN2A, anal. of methylation of; detection of nucleic acid sequences
         by isothermal RNA polymerase-dependent primer extension)
IT
      Genetic element
      RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (CpG island, anal. of methylation of; detection of nucleic acid
         sequences by isothermal RNA polymerase-dependent primer
         extension)
 IT
      Bacteriophage SP6
      Coliphage T7
      Enterobacteria phage T3
      Escherichia coli
         (RNA polymerase of; detection of nucleic acid sequences by
         isothermal RNA polymerase-dependent primer extension)
 IT
      Methylation
      RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (anal. of; detection of nucleic acid sequences by isothermal
         RNA polymerase-dependent primer extension)
 IT
      Nucleotides, analysis
      RL: ARU (Analytical role, unclassified); ANST (Analytical study)
         (analogs, as chain terminators for transcriptional primer elongation;
         detection of nucleic acid sequences by isothermal RNA
         polymerase-dependent primer extension)
IT
      Nucleoside analogs
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Nucleosides, analysis Nucleotides, analysis RL: ARU (Analytical role, unclassified); ANST (Analytical study) (as chain terminators for transcriptional primer elongation; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Fluorescent dyes (as reporter groups; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) DNA RNA RL: ANT (Analyte); ANST (Analytical study) (as template for amplification; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Diagnosis Diagnosis (cancer, anal. of DNA methylation in; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Proteins RL: ANT (Analyte); ANST (Analytical study) (conjugates with oligonucleotides, detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Primers (nucleic acid) RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Mutation (detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Pathogen (diagnostic detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Nucleic acid hybridization (for capture and anal. of amplification products; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Antibodies and Immunoglobulins RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (for protein capture; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Fluorescence resonance energy transfer (in detection of transcriptional primer extension; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Disulfides RL: MOA (Modifier or additive use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses) (in protein conjugation with oligonucleotides; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Diagnosis (mol.; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Transcription, genetic (nucleic acid amplification using; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Deamination (of 5-methylcytosine, in detection of DNA methylation; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension) Cyanine dyes

(oligonucleotide conjugates, as reporters; detection of nucleic acid

sequences by isothermal RNA polymerase-dependent primer

TI

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extension)

IT Quantum dot devices

(primer conjugates, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT Phycoerythrins

RL: ARU (Analytical role, unclassified); ANST (Analytical study) (primer conjugates, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT Promoter (genetic element)

RL: ARU (Analytical role, unclassified); BUU (Biological use, unclassified); ANST (Analytical study); BIOL (Biological study); USES (Uses)

(primers containing, for transcriptional amplification; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT mRNA

RL: ANT (Analyte); BSU (Biological study, unclassified); ANST (Analytical study); BIOL (Biological study)

(specific detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT Nucleic acid amplification (method)

(transcriptional; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 3051-11-4D, Brilliant Yellow, primer conjugates

RL: ARU (Analytical role, unclassified); ANST (Analytical study)
(Brilliant Yellow, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 18472-87-2D, Cyanosine, primer conjugates

RL: ARU (Analytical role, unclassified); ANST (Analytical study) (Cyanosine, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 146368-16-3D, Cy3, primer conjugates

RL: ARU (Analytical role, unclassified); ANST (Analytical study) (Cy3, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 146368-14-1D, primer conjugates

RL: ARU (Analytical role, unclassified); ANST (Analytical study) (Cy5, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 989-38-8D, R 6G, primer conjugates

RL: ARU (Analytical role, unclassified); ANST (Analytical study)
(R 6G, as reporter; detection of nucleic acid sequences by isothermal
RNA polymerase-dependent primer extension)

.67-43-6D, primer conjugates 81-88-9D, derivs., primer conjugates 81-88-9D, Rhodamine B, primer conjugates 83-88-5D, Riboflavin, primer 88-68-6D, Anthranilamide, primer conjugates conjugates 90-33-5D, 4-Methylumbelliferone, primer conjugates 91-64-5D, Coumarin, derivs., 129-00-0D, Pyrene, derivs., primer conjugates primer conjugates 143-74-8D, Phenol Red, primer conjugates 260-94-6D, Acridine, derivs., 569-61-9D, Pararosaniline, primer conjugates primer conjugates 574-93-6D, Phthalocyanine, primer conjugates 596-27-0D, o-Cresolphthalein, primer conjugates 605-65-2D, Dansyl chloride, primer 633-00-1D, Rosolic acid, primer conjugates conjugates 643-79-8D, o-Phthaldialdehyde, primer conjugates 2321-07-5D, Fluorescein, derivs., primer conjugates 3520-42-1D, Sulforhodamine B, primer conjugates 3546-21-2D, Ethidium, primer conjugates 3604-79-3D, m-Nitrotyrosine, 7440-27-9D, Terbium, chelates, primer conjugates primer conjugates 7612-98-8D, DABITC, primer conjugates 7613-08-3D, Acridine 2-isothiocyanate, primer conjugates 16423-68-0D, Erythrosin B, primer conjugates 16574-43-9D, Bromopyrogallol Red, primer conjugates 17372-87-1D, Eosin, derivs., primer conjugates 17681-50-4D, Reactive Red 4, primer conjugates 23627-89-6D, Naphthalocyanine, primer conjugates 25338-56-1D, Pyrenebutyric acid, primer conjugates 26093-31-2D, Coumarin 27072-45-3D, FITC, primer conjugates 120, primer conjugates

27816-59-7D, 4-Acetamido-4'-isothiocyanatostilbene-2,2'-disulfonic acid, 38183-12-9D, Fluorescamine, primer conjugates primer conjugates 47165-04-8D, DAPI, primer conjugates 50402-56-7D, EDANS, primer 51306-35-5D, DTAF, primer conjugates 53005-05-3D, 4,4'-Diisothiocyanatostilbene-2,2'-disulfonic acid, primer conjugates 53518-15-3D, 7-Amino-4-trifluoromethylcoumarin, primer conjugates 54849-69-3D, IR 144, primer conjugates 60311-02-6D, Sulforhodamine 101, primer conjugates 60520-47-0D, Eosin isothiocyanate, primer conjugates 61481-03-6D, primer conjugates 62669-70-9D, Rhodamine 123, primer 70281-37-7D, Tetramethyl rhodamine, primer conjugates conjugates 76823-03-5D, FAM, primer conjugates 82344-98-7D, XRITC, primer 82354-19-6D, Texas Red sulfonyl chloride, primer conjugates conjugates 82855-40-1D, JOE, primer conjugates 107347-53-5D, TRITC, primer 107743-39-5D, primer conjugates 120718-39-0D, ROX, primer conjugates 120718-52-7D, TAMRA, primer conjugates 138026-71-8D, conjugates BODIPY, primer conjugates 147492-82-8D, Malachite green isothiocyanate, 154088-80-9D, La Jolla Blue, primer conjugates primer conjugates 169799-14-8D, Cy7, primer conjugates 172777-84-3D, Cy5.5, primer 251102-88-2D, IRD 700, primer conjugates 256651-38-4D, IRD 800, primer conjugates 500723-56-8D, IR 1446, primer conjugates 522600-45-9D, primer conjugates 522600-44-8D, primer conjugates 524019-23-6D, primer conjugates 522600-46-0D, primer conjugates RL: ARU (Analytical role, unclassified); ANST (Analytical study) (as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 951-78-0, Deoxyuridine

RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (detection in DNA in anal. of DNA methylation; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 554-01-8, 5-Methylcytosine

RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (detection in DNA of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 9012-90-2, DNA-dependent DNA polymerase 9014-24-8, DNA-dependent RNA polymerase 9026-28-2, RNA-dependent RNA polymerase 64885-96-7, Primase RL: CAT (Catalyst use); USES (Uses)

(detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 2382-65-2D, methylated

RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (detection of; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 80057-51-8D, Erythrosin isothiocyanate, primer conjugates RL: ARU (Analytical role, unclassified); ANST (Analytical study) (erythrosin isothiocyanate, as reporter; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 144-48-9, Iodoacetamide 541-59-3, Maleimide
 RL: MOA (Modifier or additive use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)

(in protein conjugation with oligonucleotides; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

IT 524091-13-2

RL: PRP (Properties)

(unclaimed sequence; detection of nucleic acid sequences by isothermal RNA polymerase-dependent primer extension)

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TI Mechanism and stoichiometry of interaction of DnaG primase with DnaB helicase of Escherichia coli in RNA primer synthesis.

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DT Journal; Article; (JOURNAL ARTICLE) (RESEARCH SUPPORT, U.S. GOV'T, P.H.S.)

LA English

FS Priority Journals

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TI Mechanism and stoichiometry of interaction of DnaG primase with DnaB helicase of Escherichia coli in RNA primer synthesis.

Initiation and synthesis of RNA primers in the lagging strand of AB the replication fork in Escherichia coli requires the replicative DnaB helicase and the DNA primase, the DnaG gene product. In addition, the physical interaction between these two replication enzymes appears to play a role in the initiation of chromosomal DNA replication. In vitro, DnaB helicase stimulates primase to synthesize primers on single-stranded (ss) oligonucleotide templates. Earlier studies hypothesized that multiple primase molecules interact with each DnaB hexamer and single-stranded DNA. We have examined this hypothesis and determined the exact stoichiometry of primase to DnaB hexamer. We have also demonstrated that ssDNA binding activity of the DnaB helicase is necessary for directing the primase to the initiator trinucleotide and synthesis of 11-20-nucleotide long primers. Although, association of these two enzymes determines the extent and rate of synthesis of the RNA primers in vitro, direct evidence of the formation of primase-DnaB complex has remained elusive in E. coli due to the transient nature of their interaction. Therefore, we stabilized this complex. . . cross-linker and carried out a stoichiometric analysis of this complex by gel filtration. This allowed us to demonstrate that the primase-helicase complex of E. coli is comprised of three molecules of primase bound to one DnaB hexamer. Fluorescence anisotropy studies of the interaction of DnaB with primase, labeled with the fluorescent probe Ru(bipy)3, and Scatchard analysis further supported this conclusion. addition of DnaC protein, leading to the formation of the DnaB-DnaC complex, to the simple priming system resulted in the synthesis of shorter primers. Therefore, interactions of the DnaB-primase complex with other replication factors might be critical for determining the physiological length of the RNA primers in vivo and the overall kinetics of primer synthesis.

CT Anisotropy

*Bacterial Proteins

Binding Sites

Chromatography, Gel

Chromatography, High Pressure Liquid

*DNA Helicases: CH, chemistry

*DNA Helicases: ME, metabolism

*DNA Primase: CH, chemistry

*DNA Primase: ME, metabolism

*DNA Primers: CH, chemistry

DNA, Single-Stranded

DnaB Helicases

Dose-Response Relationship, Drug *Escherichia coli: EN, enzymology Escherichia coli: ME, metabolism Fluorescent Dyes Glutaral: CH, chemistry Kinetics Mutation Oligonucleotides: CH, chemistry Protein Binding *RNA: CH, chemistry 111-30-8 (Glutaral); 63231-63-0 (RNA) RN 0 (Bacterial Proteins); 0 (DNA Primers); 0 (DNA, Single-Stranded); 0 (CN Fluorescent Dyes); 0 (Oligonucleotides); EC 2.7.7.- (DNA Primase); EC 3.1.- (DnaB Helicases); EC 3.6.1.- (DNA Helicases)